the participant (or is an improved version of an existing work that the participant has sufficient rights to use and improve), and that the submission does not infringe any copyright or any other rights of any third party of which participant is aware. In addition, each participant (whether participating singly or in a group) grants to the U.S. Government a paid-up, nonexclusive, royalty-free, irrevocable worldwide license in perpetuity and the right to reproduce, publish, post, link to, share, display publicly (on the web or elsewhere) and prepare derivative works, including the right to authorize others to do so on behalf of the U.S. Government.

3. Each participant must clearly delineate any intellectual property and/or confidential commercial information contained in a submission that the participant wishes to protect as proprietary data, in accordance with Additional Rules of Participation No. 5.

4. If the submission includes any third-party works (such as third-party content or open-source code), the participant must be able to provide, upon request, documentation of all appropriate licenses and releases for use of such third-party works. If the participant cannot provide documentation of all required licenses and releases, AHRQ reserves the right, in its sole discretion, to disqualify the submission.

Marquita Cullom,

Associate Director.

[FR Doc. 2022–08908 Filed 4–26–22; 8:45 am]

BILLING CODE 4160–90–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Toxic Substances and Disease Registry

[Docket No. ATSDR–2022–0003]

Availability of Four Draft Toxicological Profiles

AGENCY: Agency for Toxic Substances and Disease Registry (ATSDR), Department of Health and Human Services (HHS).

ACTION: Notice with comment period.

SUMMARY: The Agency for Toxic Substances and Disease Registry (ATSDR), within the Department of Health and Human Services (HHS), announces the opening of a docket to obtain comments on drafts of four updated toxicological profiles: Nitrobenzene, Nitrophenols, Mercury, and Copper.

DATES: Written comments must be received on or before July 26, 2022.

ADDRESSES: You may submit comments, identified by Docket Number ATSDR–2022–0003, by either of the following methods:

- *Federal eRulemaking Portal: www.regulations.gov.* Follow the instructions for submitting comments.
- *Mail:* Office of Innovation and Analytics, Agency for Toxic Substances and Disease Registry, 4770 Buford Highway, Mail Stop S102–1, Atlanta, GA 30341–3717. Attn: Docket No. ATSDR–2022–0003.

Instructions: All submissions must include the Agency name and Docket Number. All relevant comments received will be posted without change to *http://www.regulations.gov,* including any personal information provided. Do not submit comments by email. ATSDR does not accept comments by email. For access to the docket to read background documents or comments received, go to *http://www.regulations.gov.*

FOR FURTHER INFORMATION CONTACT: Kambria Haire, Office of Innovation and Analytics, Agency for Toxic Substances and Disease Registry, 4770 Buford Highway, Mail Stop S102–1, Atlanta, GA 30329–4027; Email: *ATSDRToxProfileFRNs@cdc.gov;* Telephone: 1–800–232–4636.

SUPPLEMENTARY INFORMATION: ATSDR has prepared drafts of four updated toxicological profiles based on current understanding of the health effects and availability of new studies and other information since their initial release. All toxicological profiles issued as ''Drafts for Public Comment'' represent the result of ATSDR's evidence-based evaluations to provide important toxicological information on priority hazardous substances to the public and health professionals. ATSDR considers key studies for these substances during the profile development process. To that end, ATSDR is seeking public comments and additional information or reports on studies about the health effects of these four substances for review and potential inclusion in the profiles. ATSDR will evaluate the quality and relevance of such data or studies for possible inclusion in the profile.

Public Participation

Interested persons or organizations are invited to participate by submitting written views, information, and data. Comments received, including attachments and other supporting materials, are part of the public record and are subject to public disclosure. Comments will be posted on *https://www.regulations.gov.* Therefore, do not include any information in your comment or supporting materials that you consider confidential or inappropriate for public disclosure. If you include your name, contact information, or other information that identifies you in the body of your comments, that information will be on public display. Do not submit comments by email. ATSDR does not accept comments by email. ATSDR will review all submissions and may choose to redact or withhold submissions containing private or proprietary information such as Social Security numbers, medical information, inappropriate language, or duplicate/near duplicate examples of a mass-mail campaign. ATSDR will carefully review and consider all comments submitted in preparation of the Final Toxicological Profiles and may revise the profiles as appropriate.

Legislative Background

The Superfund Amendments and Reauthorization Act of 1986 (SARA) [42 U.S.C. 9601 *et seq.*] amended the Comprehensive Environmental Response, Compensation, and Liability Act of 1980 (CERCLA or Superfund) [42 U.S.C. 9601 *et seq.*] by establishing certain requirements for ATSDR and the U.S. Environmental Protection Agency (EPA) regarding the hazardous substances most commonly found at facilities on the CERCLA National Priorities List. Among these statutory requirements is a mandate for the Administrator of ATSDR to prepare toxicological profiles for each substance included on the priority list of hazardous substances [also called the Substance Priority List (SPL)]. This list identifies 275 hazardous substances that ATSDR and EPA have determined pose the most significant potential threat to human health. The SPL is available online at *www.atsdr.cdc.gov/spl.* ATSDR is also mandated to revise and publish updated toxicological profiles, as necessary, to reflect updated health effects and other information.

In addition, CERCLA provides ATSDR with the authority to prepare toxicological profiles for substances not found on the SPL. CERCLA authorizes ATSDR to establish and maintain an inventory of literature, research, and studies on the health effects of toxic substances (CERCLA Section 104(i)(1)(B); 42 U.S.C. 9604(i)(1)(B)); to respond to requests for health consultations (CERCLA Section 104(i)(4); 42 U.S.C. 9604(i)(4)); and to support the site-specific response actions conducted by the agency (CERCLA Section 104(i)(6); 42 U.S.C.

9604(i)(6)). Public nominations for substances from the SPL (or other substances) for toxicological profile development were requested on April 18, 2018 (83 FR 17177).

ATSDR has now prepared drafts of four updated toxicological profiles based on current understanding of the health effects and availability of new studies and other information since their initial release.

Availability

The Draft Toxicological Profiles are available online at *http://www.atsdr.cdc.gov/ToxProfiles* and at *www.regulations.gov,* Docket No. ATSDR–2022–0003.

Pamela Protzel Berman,

Associate Director, Office of Policy, Planning and Partnerships, Agency for Toxic Substances and Disease Registry.

[FR Doc. 2022–08995 Filed 4–26–22; 8:45 am]

BILLING CODE 4163–70–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Mine Safety and Health Research Advisory Committee (MSHRAC)

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice of meeting.

SUMMARY: In accordance with the Federal Advisory Committee Act, the CDC announces the following meeting for the Mine Safety and Health Research Advisory Committee (MSHRAC). This is a hybrid meeting, accessible both in-person and virtually. It is open to the public in person and limited only by the space available. Time will be available for public comment.

DATES: The meeting will be held on May 17, 2022, from 10:00 a.m. to 4:05 p.m., EDT, and May 18, 2022, from 10:00 a.m. to 4:00 p.m., EDT. The public comment session will be held on May 18, 2022, at 11:55 a.m., EDT and conclude at 12:15 p.m., EDT or following the final call for public comment, whichever comes first.

ADDRESSES: National Institute for Occupational Safety and Health (NIOSH) Pittsburgh Campus, Building 140, Room 140MP, 626 Cochrans Mill Road, Pittsburgh, Pennsylvania 15236.

Please note that the meeting location, the NIOSH Pittsburgh Campus, is a federal facility and in-person access is limited to United States citizens unless prior authorizations, taking up to 30 to 60 days, have been made. If you wish to attend either in person or virtually, please contact Ms. Berni Metzger by email at *metzger@cdc.gov* or by telephone at (412) 386–4541 at least 5 business days in advance of the meeting. If attending virtually, Ms. Metzger will provide you with the Zoom web conference access information.

Meeting Information: The conference room accommodates approximately 49 people and virtual access is limited by the number of web conference lines (500 web conference lines are available).

FOR FURTHER INFORMATION CONTACT: George W. Luxbacher, P.E., Ph.D., Designated Federal Officer, MSHRAC, NIOSH, CDC, 1600 Clifton Road, Mailstop V24–4, Atlanta, Georgia 30329–4027, Telephone: (404) 498–2808; Email: *GLuxbacher@cdc.gov.*

SUPPLEMENTARY INFORMATION:

Purpose: This committee is charged with providing advice to the Secretary, Department of Health and Human Services; the Director, CDC; and the Director, NIOSH, on priorities in mine safety and health research, including grants and contracts for such research, 30 U.S.C. 812(b)(2), Section 102(b)(2).

Matters To Be Considered: The agenda will include discussions on NIOSH mining safety and health research organizational structure, capabilities, projects, and outcomes. The meeting will also include an update from the NIOSH Associate Director for Mining. Agenda items are subject to change as priorities dictate.

Public Participation

Written Public Comment: The public may submit written comments or questions in advance of the meeting, to the contact person above. Written comments received in advance of the meeting will be included in the official record of the meeting and questions will be answered during the oral public comment period open to public participation.

Oral Public Comment: This meeting will include time for members of the public to make an oral comment. The public comment session will be held on May 18, 2022, at 11:55 a.m., EDT and conclude at 12:15 p.m., EDT or following the final call for public comment, whichever comes first.

The Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Kalwant Smagh,

Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention.

[FR Doc. 2022–08926 Filed 4–26–22; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Community Living

Agency Information Collection Activities; Submission for OMB Review; Public Comment Request; of the No Wrong Door (NWD) System Management Tool OMB Control 0985–0062

AGENCY: Administration for Community Living, HHS.

ACTION: Notice.

SUMMARY: The Administration for Community Living is announcing that the proposed collection of information listed above has been submitted to the Office of Management and Budget (OMB) for review and clearance as required under section 506(c)(2)(A) of the Paperwork Reduction Act of 1995. This 30-Day notice collects comments on the information collection requirements related to the information collection requirements of the No Wrong Door (NWD) System Management Tool OMB Control 0985–0062.

DATES: Submit written comments on the collection of information by May 27, 2022.

ADDRESSES: Submit written comments and recommendations for the proposed information collection within 30 days of publication of this notice to *www.reginfo.gov/public/do/PRAMain.* Find the information collection by selecting ''Currently under 30-day Review—Open for Public Comments'' or by using the search function. By mail to the Office of Information and Regulatory Affairs, OMB, New Executive Office Bldg., 725 17th St. NW, Rm. 10235, Washington, DC 20503, Attn: OMB Desk Officer for ACL.

FOR FURTHER INFORMATION CONTACT: Kristie Kulinski, (202) 795–7379 or *kristie.kulinski@acl.hhs.gov.* Administration for Community Living.

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, ACL has submitted the following proposed collection of information to OMB for review and clearance.